10

1. A method for vaccinating a human against a human immunodeficiency virus comprising the steps of:

selecting an immunogen competent to induce a protective immune response in said human against said human immunodeficiency virus, and

administering to said human an effective amount of said immunogen sufficient to induce a sustained cell mediated immune response against said human immunodeficiency virus.

- 2. The method of claim 1 wherein said immunogen is an attenuated form of human immunodeficiency virus.
- 3. The method of claim 2/wherein said immunogen has been attenuated by removing all or part of the nef gene from the nucleic acid of said human immunodeficiency virus.
- 4. The method of claim 1 wherein said immunogen is a subunit of said human immunodeficiency virus.
- 5. The method of claim 4 wherein said immunogen is a gp120 subunit of said human immunodeficiency virus.
- 6. The method of claim 4 wherein said immunogen is a gp160 subunit of said human immunodeficiency virus.
- 7. The method of claim 1 wherein said immunogen is an inactivated human immunodeficiency virus.
- 8. The method of claim 7 wherein said immunogen has been inactivated by removing a sufficient portion of its genetic material so as to render it incapable of replicating.
- 9. The method of claim 8 wherein the genetic material removed from said human immunodeficiency virus is a portion of a gene coding for a gag nucleocapsid protein.

35

25

30

- 11. The method of claim 1 wherein said immunogen is an infectious form of human immunodeficiency virus administered in a subinfectious amount.
- 12. The method of claim 1 wherein the effective amount of immunogen administered contains between 100 attograms and 20 milligrams of p24 gag antigen.
- 13. The method of claim 2 wherein the effective amount of immunogen administered contains between 10 and 500 femtograms of p24 gag antigen.
- 14. The method of claim 11 wherein the effective amount of immunogen administered contains between 100 attograms and 500 femtograms of p24 gag antigen.
- 15. The method of claim 1 wherein a cell mediated response is determined to be present using a T-Cell proliferation assay if the uptake of thymidine by antigenstimulated cells is at least four-fold above background.
- response is determined to be present using an IL-2 assay if the production of IL-2 by antigen-stimulated cells is at least four-fold above background.
- 17. A method for vaccinating a human against a human immunodeficiency virus comprising the steps of:

 selecting an immunogen competent to induce a protective immune response in said human against said human immunodeficiency virus, and

administering an effective amount of said immunogen to said human sufficient to induce a cell mediated

09769NN3 012401

5

10

25

30

35

response against said human immunodeficiency virus but below the amount necessary to induce a humoral response.

18. A method for vaccinating a human against a mammalian retrovirus comprising the steps of:

selecting an immunogen competent to induce a protective immune response in said mammal against said retrovirus, and

administering an effective amount of said immunogen to said mammal sufficient to induce a cell mediated immune response against/said/retrovirus but below the level necessary to induce /a wurdral response.

The method of claim 18 wherein said retrovirus is a simian immunodeficiency virus.

human

The method of claim 18 wherein said mammal is a

The method of claim 20 wherein said retrovirus 21. is HTLV-I.

The method of claim 20 wherein said retrovirus is HTLV-II.

25

- 23. The method of claim 20 wherein said retrovirus is foamy virus.
- A vaccine comprising a therapeutically 30 effective dose of an /mmungen/capable of eliciting a cellmediated immune/response in a human protective against infection by a/human immunodeliciency virus.
- A/vaccine comprising a dose of immunogen 35 capable of elicating a cell-mediated response in a human as measured by a /r-cell proliferation assay.

5

10